EAST Search History

Ref #	Hits	Search Query	DBs	Default Operator	Plurals	Time Stamp
L1	93	(540/462,549/270,560/51,568/449). CCLS.	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	OFF	2007/08/13 12:31
L2	19	L1 AND EPOTHILONE	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	OFF	2007/08/13 12:31

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        MAY 08
                 CA/CAplus Indian patent publication number format defined
NEWS
        MAY 14
                 RDISCLOSURE on STN Easy enhanced with new search and display
                 fields
NEWS
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        MAY 21
                 BIOSIS reloaded and enhanced with archival data
NEWS
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        MAY 21
                TOXCENTER enhanced with BIOSIS reload
NEWS
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        MAY 21
                CA/CAplus enhanced with additional kind codes for German
                 patents
        MAY 22
NEWS
                CA/CAplus enhanced with IPC reclassification in Japanese
                 patents
                CA/CAplus enhanced with pre-1967 CAS Registry Numbers
        JUN 27
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NEWS 10
        JUN 29
                 STN Viewer now available
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NEWS 11
                STN Express, Version 8.2, now available
NEWS 12
        JUL 02
                LEMBASE coverage updated
NEWS 13 JUL 02 LMEDLINE coverage updated
NEWS 14 JUL 02
                SCISEARCH enhanced with complete author names
NEWS 15 JUL 02
                CHEMCATS accession numbers revised
NEWS 16 JUL 02 CA/CAplus enhanced with utility model patents from China
        JUL 16 CAplus enhanced with French and German abstracts
NEWS 17
NEWS 18
        JUL 18
                CA/CAplus patent coverage enhanced
NEWS 19
        JUL 26 USPATFULL/USPAT2 enhanced with IPC reclassification
NEWS 20 JUL 30
                USGENE now available on STN
NEWS 21
        AUG 06
                CAS REGISTRY enhanced with new experimental property tags
NEWS 22'
        AUG 06
                BEILSTEIN updated with new compounds
NEWS 23
        AUG 06
                FSTA enhanced with new thesaurus edition
NEWS 24
        AUG 13
                CA/CAplus enhanced with additional kind codes for granted
                 patents
NEWS EXPRESS
             29 JUNE 2007: CURRENT WINDOWS VERSION IS V8.2,
              CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
              AND CURRENT DISCOVER FILE IS DATED 05 JULY 2007.
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FULL ESTIMATED COST

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chain nodes :
4 7 11 12 14 16 23 24 26 27 28 29 30 31
                                                                  32
                                                                      33 34 35 36 37 39
40 41
ring nodes :
1 2 3 5 6 8 9 10 13 15 17 18
                                                  19 20 21 25
chain bonds :
1-4 \quad 3-23 \quad 5-28 \quad 5-29 \quad 6-7 \quad 8-39 \quad 9-11 \quad 10-12 \quad 10-40 \quad 13-14 \quad 15-16 \quad 15-41 \quad 17-30
17-31 18-32 18-33 19-34 19-35 23-24 23-37 24-36 25-26 25-27
ring bonds :
1-2 1-5 2-3 3-25 5-6 6-8 8-9 9-10 10-13 13-15 15-17 17-18
                                                                                     18-19 19-20
20-21 21-25
exact/norm bonds :
1-2 1-4 1-5 2-3 3-23 3-25 5-6 5-28 5-29 6-7 6-8 8-9 8-39 9-10 9-11 10-12 10-13 10-40 13-14 13-15 15-16 15-17 15-41 17-18 17-30 17-31 18-19 18-32 18-33 19-20 19-34 19-35 20-21 21-25 23-24 23-37 24-36 25-26 25-27
isolated ring systems :
containing 1 :
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Match level:

1:Atom 2:Atom 3:Atom 4:CLASS 5:Atom 6:Atom 7:CLASS 8:Atom 9:Atom 10:Atom 11:CLASS 12:CLASS 13:Atom 14:CLASS 15:Atom 16:CLASS 17:Atom 18:Atom 19:Atom 20:Atom 21:Atom 23:CLASS 24:CLASS 25:Atom 26:CLASS 27:CLASS 28:CLASS 29:CLASS 30:CLASS 31:CLASS 32:CLASS 33:CLASS 34:CLASS 35:CLASS 36:CLASS 37:CLASS 39:CLASS 40:CLASS 41:CLASS

L1 STRUCTURE UPLOADED

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chain nodes : 4 7 11 12 14 16 23 24 26 27 28 29 30 31 32 33 34 35 36 37 39 40 41 ring nodes : 1 2 3 5 6 8 9 10 13 15 17 18 19 20 21 25 42 chain bonds : 1-4 3-23 5-28 5-29 6-7 8-39 9-11 10-12 10-40 13-14 15-16 15-41 17-30 17-31 18-32 18-33 19-34 19-35 23-24 23-37 24-36 25-26 25-27

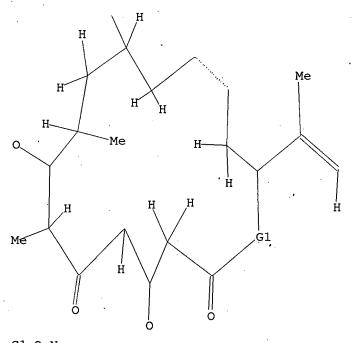
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Match level:

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L2 STRUCTURE UPLOADED

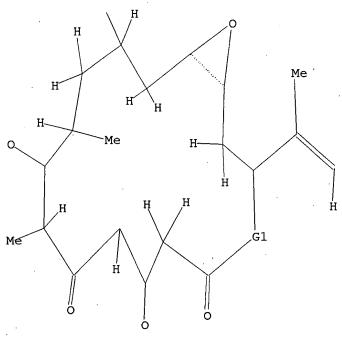
=> d l1 L1 HAS NO ANSWERS L1 STR



G1 O,N

Structure attributes must be viewed using STN Express query preparation.

=> d 12 L2 HAS NO ANSWERS L2 STR



G1 O, N

Structure attributes must be viewed using STN Express query preparation.

=> s l1 full

FULL SEARCH INITIATED 10:20:46 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 56309 TO ITERATE

100.0% PROCESSED 56309 ITERATIONS SEARCH TIME: 00.00.01

14 ANSWERS

L3 ·

14 SEA SSS FUL L1

=> s 12 full

FULL SEARCH INITIATED 10:20:52 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 1862 TO ITERATE

100.0% PROCESSED 1862 ITERATIONS

9 ANSWERS

SEARCH TIME: 00.00.01

114

9 SEA SSS FUL L2

=> file caplus

COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 344.65 344.86

FULL ESTIMATED COST

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=> s 13 full

L5 3 L3

=> d ibib abs hitstr tot

L5 ANSWER 1 OF 3 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

2005:460208 CAPLUS

DOCUMENT NUMBER:

143:171398

TITLE:

Production of epothilones derivatives in Myxococcus or

Sorangium comprising PKS mutant gene

INVENTOR(S):

Qiu, Rongquo

PATENT ASSIGNEE(S):

Beijing Huahao Zhongtian Biotechnology Co., Ltd.,

Peop. Rep. China

SOURCE:

Faming Zhuanli Shenqing Gongkai Shuomingshu, No pp.

given

CODEN: CNXXEV

DOCUMENT TYPE:

Patent

LANGUAGE:

Chinese

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
	-			
CN 1521258	Α	20040818	CN 2003-103031,	20030128
PRIORITY APPLN. INFO.:			CN 2003-103031	20030128
OTHER SOURCE(S):	CASRE	ACT 143:1713	398; MARPAT 143:171398	
			of epothilones derivs.	in Myxococcus or
Sorangium comprisi	ng PKS i	mutant gene.	The invention also r	elates to the
uses of these comp	ds. in	preparing me	edicine composition for	treating tumor.
inhibiting cell pr	olifera	tion and gro	owth.	ordering camer,
IT 252917-35-4P 25291				

RL: BPN (Biosynthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(production of epothilones derivs. in Myxococcus or Sorangium comprising PKS mutant gene)

RN 252917-35-4 CAPLUS

CN Oxacyclohexadec-13-ene-2,6-dione, 4,8-dihydroxy-5,7,9,13-tetramethyl-16-[(1E)-1-methyl-2-(2-methyl-4-thiazolyl)ethenyl]-, (4S,7R,8S,9S,13Z,16S)-(9CI) (CA INDEX NAME)

RN 252917-37-6 CAPLUS

CN Oxacyclohexadec-13-ene-2,6-dione, 4,8-dihydroxy-5,7,9,13-tetramethyl-16-[(1E)-1-methyl-2-(2-methyl-4-thiazolyl)ethenyl]-, (4S,7R,8S,9S,13Z,16S)-(9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (-). Double bond geometry as shown. Currently available stereo shown.

RN 860300-23-8 CAPLUS

CN Oxacyclohexadec-13-ene-2,6-dione, 4,8-dihydroxy-16-[(1E)-2-[2-(hydroxymethyl)-4-thiazolyl]-1-methylethenyl]-5,7,9,13-tetramethyl-, (4S,7R,8S,9S,13Z,16S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Double bond geometry as shown.

IT 860300-09-0P 860300-14-7P 860300-16-9P 860300-17-0P 860300-18-1P 860300-20-5P 860300-26-1P

RL: SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(production of epothilones derivs. in Myxococcus or Sorangium comprising PKS mutant gene)

RN 860300-09-0 CAPLUS

CN Oxacyclohexadec-13-ene-2,6-dione, 4-hydroxy-8-methoxy-5,7,9,13-tetramethyl-16-[(1E)-1-methyl-2-(2-methyl-4-thiazolyl)ethenyl]-, (4S,7R,8S,9S,13Z,16S)-(9CI) (CA INDEX NAME)

Absolute stereochemistry.

Double bond geometry as shown.

RN 860300-14-7 CAPLUS

CN Oxacyclohexadecane-2,6-dione, 4,13-dihydroxy-8-methoxy-5,7,9,13-tetramethyl-16-[(1E)-1-methyl-2-(2-methyl-4-thiazolyl)ethenyl]-, (4S,7R,8S,9S,16S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.
Double bond geometry as shown.

RN 860300-16-9 CAPLUS

CN Oxacyclohexadecane-2,6-dione, 4,8,13-trihydroxy-5,7,9,13-tetramethyl-16-[(1E)-1-methyl-2-(2-methyl-4-thiazolyl)ethenyl]-, (4S,7R,8S,9S,16S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

Double bond geometry as shown.

RN 860300-17-0 CAPLUS

CN Oxacyclohexadecane-2,6-dione, 4,8,13-trihydroxy-5,7,9,13-tetramethyl-16-[(1E)-1-methyl-2-(2-methyl-4-oxazolyl)ethenyl]-, (4S,7R,8S,9S,16S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Double bond geometry as shown.

RN 860300-18-1 CAPLUS

CN Azacyclohexadec-13-ene-2,6-dione, 4,8-dihydroxy-5,7,9,13-tetramethyl-16-[(1E)-1-methyl-2-(2-methyl-4-thiazolyl)ethenyl]-, (4S,7R,8S,9S,13Z,16S)-(9CI) (CA INDEX NAME)

Absolute stereochemistry. Double bond geometry as shown.

RN 860300-20-5 CAPLUS

CN Oxacyclohexadec-13-ene-2,6-dione, 16-[(1E)-2-[2-(aminomethyl)-4-thiazolyl]-1-methylethenyl]-4,8-dihydroxy-5,7,9,13-tetramethyl-, (4S,7R,8S,9S,13Z,16S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Double bond geometry as shown.

RN 860300-26-1 CAPLUS

CN Oxacyclohexadec-13-ene-2,6-dione, 4,8-dihydroxy-5,7,9,13-tetramethyl-16-[(1E)-1-methyl-2-(2-methyl-4-oxazolyl)ethenyl]-, (4S,7R,8S,9S,13Z,16S)-(9CI) (CA INDEX NAME)

Absolute stereochemistry.
Double bond geometry as shown.

L5 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2001:413810 CAPLUS

DOCUMENT NUMBER: 135:179755

TITLE: New Natural Epothilones from Sorangium cellulosum,

Strains So ce90/B2 and So ce90/D13: Isolation,

Structure Elucidation, and SAR Studies

AUTHOR(S): Hardt, Ingo H.; Steinmetz, Heinrich; Gerth, Klaus;

Sasse, F.; Reichenbach, Hans; Hoefle, Gerhard

CORPORATE SOURCE: Gesellschaft fuer Biotechnologische Forschung mbH,

Braunschweig, D-38124, Germany

SOURCE: Journal of Natural Products (2001), 64(7), 847-856

CODEN: JNPRDF; ISSN: 0163-3864

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal LANGUAGE: English

AB In addition to epothilones A (1) and B (2), 37 natural epothilone variants

and epothilone-related compds. were isolated from the culture broth of a 700 L fermentation of Sorangium cellulosum, strain So ce90/B2. Of these, only the 12,13-desoxyepothilones, epothilone C (14) and D (15), were produced in significant amts. (3-6 mg/L); the 21-hydroxy derivs. and epothilones E (3) and F (4), in low and variable amts. due to further degradation by the producing organism. Most of the other epothilone variants were produced only in 1-100 μ g/L amts. The new compds. are very similar in structure to the parent compds. 1, 2 and 14, 15 and are presumably the result of the imperfect selectivity of the biosynthetic enzymes for acetate and propionate. Further, epothilones containing an oxazole moiety (10-13) in the side chain instead of a thiazole as well as ring-expanded 18-membered macrolides, epothilones I (30-35), and a ring contracted 14-membered macrolide, epothilone K (36), were found as very minor metabolites. mutant strain, So ce90/D13, instead of macrolactones, produced short-chain carboxylic acids 40, 41, and 42 bearing the characteristic thiazole side chain. The structures of the new epothilones were elucidated on the basis of comprehensive NMR and MS data. The new epothilone variants were tested in a cytotoxicity assay with mouse fibroblasts (cell line L929), and structure-activity relationships were established. Several new natural epothilones showed activity comparable to 1 and 2, but in no case exceeded that of 2.

IT 252917-34-3P, Epothilone C1 252917-35-4P, Epothilone D1 252917-36-5P, Epothilone C2 252917-37-6P, Epothilone D2 252917-48-9P, trans-Epothilone C1 252917-49-0P, trans-Epothilone C2

RL: BPN (Biosynthetic preparation); PRP (Properties); PUR (Purification or recovery); BIOL (Biological study); PREP (Preparation) (new natural epothilones from Sorangium cellulosum)

RN 252917-34-3 CAPLUS

CN Oxacyclohexadec-13-ene-2,6-dione, 4,8-dihydroxy-5,7,9-trimethyl-16-[(1E)-1-methyl-2-(2-methyl-4-thiazolyl)ethenyl]-, (4S,7R,8S,9S,13Z,16S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (-). Double bond geometry as shown. Currently available stereo shown.

RN 252917-35-4 CAPLUS

CN Oxacyclohexadec-13-ene-2,6-dione, 4,8-dihydroxy-5,7,9,13-tetramethyl-16-[(1E)-1-methyl-2-(2-methyl-4-thiazolyl)ethenyl]-, (4S,7R,8S,9S,13Z,16S)-(9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (-). Double bond geometry as shown.

Currently available stereo shown.

RN 252917-36-5 CAPLUS

CN Oxacyclohexadec-13-ene-2,6-dione, 4,8-dihydroxy-5,7,9-trimethyl-16-[(1E)-1-methyl-2-(2-methyl-4-thiazolyl)ethenyl]-, (4S,7R,8S,9S,13Z,16S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (-). Double bond geometry as shown. Currently available stereo shown.

RN 252917-37-6 CAPLUS

CN Oxacyclohexadec-13-ene-2,6-dione, 4,8-dihydroxy-5,7,9,13-tetramethyl-16-[(1E)-1-methyl-2-(2-methyl-4-thiazolyl)ethenyl]-, (4S,7R,8S,9S,13Z,16S)-(9CI) (CA INDEX NAME)

RN 252917-48-9 CAPLUS

CN Oxacyclohexadec-13-ene-2,6-dione, 4,8-dihydroxy-5,7,9-trimethyl-16-[(1E)-1-methyl-2-(2-methyl-4-thiazolyl)ethenyl]-, (4S,7R,8S,9S,13E,16S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (-). Double bond geometry as shown. Currently available stereo shown.

RN 252917-49-0 CAPLUS

CN Oxacyclohexadec-13-ene-2,6-dione, 4,8-dihydroxy-5,7,9-trimethyl-16-[(1E)-1-methyl-2-(2-methyl-4-thiazolyl)ethenyl]-, (4S,7R,8S,9S,13E,16S)- (9CI) (CA INDEX NAME)

REFERENCE COUNT:

THERE ARE 45 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

1999:811249 CAPLUS

DOCUMENT NUMBER:

132:49105

TITLE:

Epothilone minor constituents

INVENTOR(S):

Hoefle, Gerhard; Reichenbach, Hans; Gerth, Klaus; Hardt, Ingo; Sasse, Florenz; Steinmetz, Heinrich

PATENT ASSIGNEE(S):

Gesellschaft Fur Biotechnologische Forschung m.b.H.

(Gbf), Germany

SOURCE:

PCT Int. Appl., 36 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

German 1

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PA	PATENT NO.					ND DATE				APPL	ICAT		DATE					
WO	9965	913			A2		1999	 1223	,	WO 1	999-		19990618					
WO.	9965																	
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		ES,	FΙ,	FR,	GB,	GR,	ΙE,	IT,	LU,	MC,	NL,	PT,	SE,	BF,	ВJ,	CF,	CG,	
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ΑU	J 9948995				Α		2000	0105		AU 1	999-	4899	5		1	9990		
ΑU	J 757452				· B2		2003	0220										
EP	1087	975			A 2		2001	0404		EP 1	999-	9327	00		1	9990	618	
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JP	2002	5183	97		T .	•	2002	0625		JP 2	000-		19990618					
ΕP	1275	648			A1		2003	0115	EP 2002-22332						19990618			
	R:	ΑT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	IT,	LI,	LU,	NL,	SE.	MC.	PT.	
		ΙE,	·SI,	LT,	LV,	FI,	RO,	MK,	CY,	\mathtt{AL}						,	,	
ΑT	2481	74			T		2003	0915	AT 1999-932700 PT 1999-932700						1	9990	618	
PT	1087	975			T		2004	0130]	PT 1	999-	9327	00		1	0000	610	
ES	2207	249			Т3		2004	0516]	ES 1	999-	9327	00		1	9990	618	
US	6624	310	•		В1		2003	0923	ES 1999-932700 US 2001-719932						20010321			
US	2004	0490	51		A1		2004	0311	Ţ	US 2	003-	4570	98		20030606			
US	2006	1425							US 2003-457098 US 2006-354769						20060215			

PRIORITY APPLN. INFO.:

DE 1998-19826988 A 19980618 EP 1999-932700 A3 19990618 WO 1999-EP4244 W 19990618 US 2001-719932 A3 20010321 US 2003-457098 A1 20030606

AB The invention relates to compds. which are obtained by fermenting DSM 6773, especially epothilones A1, A2, A8, A9, B10, C1, C2, C3, C4, C5, C6, C7, C8, C9, D1, D2, D5, G1, G2, H1, H2, I1, I2, I3, I4, I5, I6 and K and trans-epothilones C1 and C2.

IT 252917-34-3P, Epothilone C1 252917-35-4P, Epothilone D1
252917-36-5P, Epothilone C2 252917-37-6P, Epothilone D2
252917-48-9P, trans-Epothilone C1 252917-49-0P,
trans-Epothilone C2

В2

RL: BAC (Biological activity or effector, except adverse); BOC (Biological occurrence); BSU (Biological study, unclassified); PRP (Properties); PUR (Purification or recovery); BIOL (Biological study); OCCU (Occurrence); PREP (Preparation)

(epothilone minor constituents)

RN 252917-34-3 CAPLUS

CN Oxacyclohexadec-13-ene-2,6-dione, 4,8-dihydroxy-5,7,9-trimethyl-16-[(1E)-1-methyl-2-(2-methyl-4-thiazolyl)ethenyl]-, (4S,7R,8S,9S,13Z,16S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (-). Double bond geometry as shown. Currently available stereo shown.

RN 252917-35-4 CAPLUS

CN Oxacyclohexadec-13-ene-2,6-dione, 4,8-dihydroxy-5,7,9,13-tetramethyl-16-[(1E)-1-methyl-2-(2-methyl-4-thiazolyl)ethenyl]-, (4S,7R,8S,9S,13Z,16S)-(9CI) (CA INDEX NAME)

RN 252917-36-5 CAPLUS

CN Oxacyclohexadec-13-ene-2,6-dione, 4,8-dihydroxy-5,7,9-trimethyl-16-[(1E)-1-methyl-2-(2-methyl-4-thiazolyl)ethenyl]-, (4S,7R,8S,9S,13Z,16S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (-). Double bond geometry as shown. Currently available stereo shown.

RN 252917-37-6 CAPLUS

CN Oxacyclohexadec-13-ene-2,6-dione, 4,8-dihydroxy-5,7,9,13-tetramethyl-16-[(1E)-1-methyl-2-(2-methyl-4-thiazolyl)ethenyl]-, (4S,7R,8S,9S,13Z,16S)-(9CI) (CA INDEX NAME)

RN 252917-48-9 CAPLUS

CN Oxacyclohexadec-13-ene-2,6-dione, 4,8-dihydroxy-5,7,9-trimethyl-16-[(1E)-1-methyl-2-(2-methyl-4-thiazolyl)ethenyl]-, (4S,7R,8S,9S,13E,16S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (-). Double bond geometry as shown. Currently available stereo shown.

RN 252917-49-0 CAPLUS

CN Oxacyclohexadec-13-ene-2,6-dione, 4,8-dihydroxy-5,7,9-trimethyl-16-[(1E)-1-methyl-2-(2-methyl-4-thiazolyl)ethenyl]-, (4S,7R,8S,9S,13E,16S)- (9CI) (CA INDEX NAME)

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L6 ANSWER 1 OF 6 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

2005:460208 CAPLUS

DOCUMENT NUMBER:

143:171398

TITLE:

Production of epothilones derivatives in Myxococcus or

Sorangium comprising PKS mutant gene

INVENTOR(S):

Qiu, Rongguo

PATENT ASSIGNEE(S):

Beijing Huahao Zhongtian Biotechnology Co., Ltd.,

Peop. Rep. China

SOURCE:

Faming Zhuanli Shenqing Gongkai Shuomingshu, No pp.

given

CODEN: CNXXEV

DOCUMENT TYPE:

P

TANGUAGE

Patent

LANGUAGE:

Chinese

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
CN 1521258	Α	20040818	CN 2003-103031	20030128
PRIORITY APPLN. INFO.:			CN 2003-103031	20030128
OTHER SOURCE(S):	CASREA	ACT 143:1713	98; MARPAT 143:171398	

AB Described is a method for production of epothilones derivs. in Myxococcus or Sorangium comprising PKS mutant gene. The invention also relates to the uses of these compds. in preparing medicine composition for treating tumor, inhibiting cell proliferation and growth.

IT 502619-65-0P 860300-22-7P

RL: BPN (Biosynthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(production of epothilones derivs. in Myxococcus or Sorangium comprising PKS mutant gene)

RN 502619-65-0 CAPLUS

CN 4,17-Dioxabicyclo[14.1.0]heptadecane-5,9-dione, 7,11-dihydroxy-8,10,12,16-tetramethyl-3-[(1E)-1-methyl-2-(2-methyl-4-thiazolyl)ethenyl]-, (1S,3S,7S,10R,11S,12S,16R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.
Double bond geometry as shown.

RN 860300-22-7 CAPLUS

CN 4,17-Dioxabicyclo[14.1.0]heptadecane-5,9-dione, 7,11-dihydroxy-3-[(1E)-2-[2-(hydroxymethyl)-4-thiazolyl]-1-methylethenyl]-8,10,12,16-tetramethyl-, (1S,3S,7S,10R,11S,12S,16R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

Double bond geometry as shown.

IT 860300-10-3P 860300-19-2P 860300-21-6P 860300-25-0P

RL: SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(production of epothilones derivs. in Myxococcus or Sorangium comprising PKS mutant gene)

RN 860300-10-3 CAPLUS

CN 4,17-Dioxabicyclo[14.1.0]heptadecane-5,9-dione, 7-hydroxy-11-methoxy-8,10,12,16-tetramethyl-3-[(1E)-1-methyl-2-(2-methyl-4-thiazolyl)ethenyl]-, (1R,3S,7S,10R,11S,12S,16S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

Double bond geometry as shown.

RN 860300-19-2 CAPLUS

CN 17-Oxa-4-azabicyclo[14.1.0]heptadecane-5,9-dione, 7,11-dihydroxy-8,10,12,16-tetramethyl-3-[(1E)-1-methyl-2-(2-methyl-4-thiazolyl)ethenyl]-, (1R,3S,7S,10R,11S,12S,16S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

Double bond geometry as shown.

RN 860300-21-6 CAPLUS

CN 4,17-Dioxabicyclo[14.1.0]heptadecane-5,9-dione, 3-[(1E)-2-[2-(aminomethyl)-4-thiazolyl]-1-methylethenyl]-7,11-dihydroxy-8,10,12,16-tetramethyl-, (1S,3S,7S,10R,11S,12S,16R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

Double bond geometry as shown.

RN 860300-25-0 CAPLUS

CN 4,17-Dioxabicyclo[14.1.0]heptadecane-5,9-dione, 7,11-dihydroxy-8,10,12,16-tetramethyl-3-[(1E)-1-methyl-2-(2-methyl-4-oxazolyl)ethenyl]-, (1S,3S,7S,10R,11S,12S,16R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Double bond geometry as shown.

ANSWER 2 OF 6 CAPLUS COPYRIGHT 2007 ACS on STN

2004:550960 CAPLUS ACCESSION NUMBER:

DOCUMENT NUMBER:

141:106321

TITLE:

Preparation of epothilone derivatives for use in pharmaceutical compositions as antitumor agents Denni-Dischert, Donatienne; Floersheimer, Andreas;

INVENTOR(S):

Kuesters, Ernst; Oberer, Lukas; Sedelmeier, Gottfried

Novartis A.-G., Switz.; Novartis Pharma G.m.b.H.

PATENT ASSIGNEE(S):

PCT Int. Appl., 50 pp.

SOURCE:

CODEN: PIXXD2

DOCUMENT TYPE:

Patent English

LANGUAGE:

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PA	TENT	NO.			KIN	D .	DATE		APPLICATION NO.							DATE			
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EP	1581	536			Δ2		2005	1005	EP 2003-785920						20031222				
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DR CN	2003	170	93		A		2005	1122	•	BR Z	003-	1/69.	3		2	0031	222		
CN	1732 2006	1/2			A		2006	0208	- (CN 2	003-	BOTO.	/416		2	0031	222		
JP	2006	5140	25		Т		2006	0427	•	JP 2	004-	5614	16		2				
	US 2006014796						2006	0119								0050			
PRIORIT	PRIORITY APPLN. INFO.:										002-					0021	223		
							•		I	WO 2	003-1	EP14	747	7	W 2	0031	222		
OTHER S	OTHER SOURCE(S):						141:	10632	321										
GI																			

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

C4-demethyl-epothilones or C4-bisnor-epothilones, such as I [R1, R7 = H, alkyl; R2 = nitrogen containing heteroaryl; R3 = H, Me; X = O, NR7; Z = O, bond], were prepared via fermentation and organic synthesis for use in pharmaceutical

compns. as antitumor agents. Thus, C4-bisnor-epothilone B II (R3 = H) was prepared via an aldol condensation of aldehyde III with in situ disilylated (3R)-3-hydroxy-5-oxoheptanoic acid followed by a desilylation/macrolactonization reaction sequence. Also, C4-demethyl-epothilone B II (R = Me) was prepared directly by a fermentation process. The prepared epothilones were assayed for efficacy against human KB-31 and KB-8511 carcinoma cells. Drug delivery formulations containing the prepared epithilones were presented.

ΙT 502619-65-0P

> RL: BPN (Biosynthetic preparation); PAC (Pharmacological activity); PUR (Purification or recovery); THU (Therapeutic use); BIOL (Biological

study); PREP (Preparation); USES (Uses)

(preparation of epothilone derivs. via fermentation and organic synthesis for use in

pharmaceutical compns. as antitumor agents)

RN 502619-65-0 CAPLUS

CN 4,17-Dioxabicyclo[14.1.0]heptadecane-5,9-dione, 7,11-dihydroxy-8,10,12,16-tetramethyl-3-[(1E)-1-methyl-2-(2-methyl-4-thiazolyl)ethenyl]-, (1S,3S,7S,10R,11S,12S,16R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

Double bond geometry as shown.

IT 717917-47-0P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

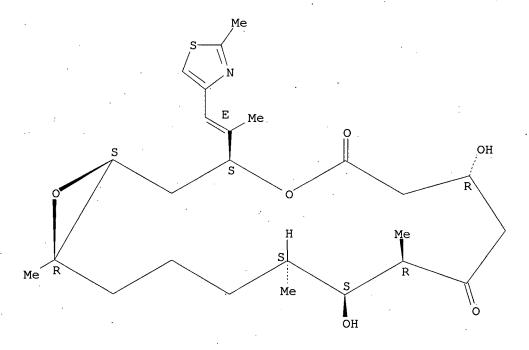
(preparation of epothilone derivs. via fermentation and organic synthesis for use in

pharmaceutical compns. as antitumor agents)

RN 717917-47-0 CAPLUS

CN 4,17-Dioxabicyclo[14.1.0]heptadecane-5,9-dione, 7,11-dihydroxy-10,12,16-trimethyl-3-[(1E)-1-methyl-2-(2-methyl-4-thiazolyl)ethenyl]-, (1S,3S,7R,10R,11S,12S,16R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.
Double bond geometry as shown.



ANSWER 3 OF 6 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

2004:182886 CAPLUS

DOCUMENT NUMBER:

140:217439

TITLE:

Synthesis of epothilones for use in pharmaceutical

compositions for the treatment of cancer

INVENTOR(S):

Danishefsky, Samuel J.; Rivkin, Alexey; Yoshimura, Fumihiko; Gabarda Ortega, Ana Esther; Cho, Young Shin;

Chou, Ting-Chao; Dongm, Huajin

PATENT ASSIGNEE(S):

Sloan-Kettering Institute for Cancer Research, USA

SOURCE:

PCT Int. Appl., 223 pp. CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE: 3

English

PATENT INFORMATION:

FAMILY ACC. NUM. COUNT:

PA'	PATENT NO.					KIND DATE		•	APPLICATION NO.						DATE			
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EP 1506203					B1		2007								20030022			

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AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK
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                                                                      20050318
PRIORITY APPLN. INFO.:
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                                                                     .20021101
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                                                                      20030320
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                                                                   Ρ
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                                              WO 2003-US26367
                                                                   W
                                                                      20030822
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Ι

OTHER SOURCE(S):

CASREACT 140:217439; MARPAT 140:217439

GI

AΒ Epothilones, such as I [R = Me, CH2OH, CH2NH2, etc.; R1 = H, Me, CF3, etc.; X = 0, bond; 9,10-saturated or -unsatd.], were prepared for therapeutic use as antitumor agents. Thus, II was prepared via a multistep synthetic sequence which included an intramol. metathesis reaction to form the macrocyclic ring. The prepared epothilones were assayed for pharmacol. activity by various means which included growth inhibition of CCRF-CEM cells.

IT 502619-65-0

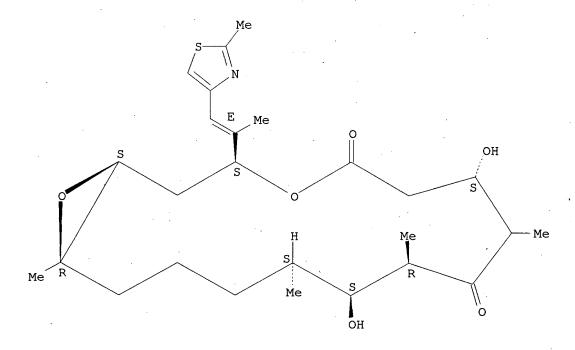
> RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(synthesis of epothilones for use in pharmaceutical compns. for the treatment of cancer)

RN 502619-65-0 CAPLUS

CN 4,17-Dioxabicyclo[14.1.0]heptadecane-5,9-dione, 7,11-dihydroxy-8,10,12,16tetramethyl-3-[(1E)-1-methyl-2-(2-methyl-4-thiazolyl)ethenyl]-, (1S, 3S, 7S, 10R, 11S, 12S, 16R) - (9CI) (CA INDEX NAME)

Absolute stereochemistry. Double bond geometry as shown.



ANSWER 4 OF 6 CAPLUS COPYRIGHT 2007 ACS on STN

138:255008

ACCESSION NUMBER:

2003:221685 CAPLUS

DOCUMENT NUMBER: TITLE:

Synthesis of epothilones for therapeutic use as

anticancer agents

INVENTOR(S):

Danishefsky, Samuel J.; Biswas, Kaustav; Chapell, Mark; Lin, Hong; Njardarson, Jon T.; Lee, Chulbom;

Rivkin, Alexey; Chou, Ting-Chao

PATENT ASSIGNEE(S):

Sloan-Kettering Institute for Cancer Research, USA

PCT Int. Appl., 219 pp.

SOURCE:

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PA	PATENT NO.					KIND DATE				APPL	ICAT:		DATE				
					A2 20030320			1	WO 2	002-1		20020906					
	2003						2004										
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								1	US 2	001-	3515	76P			0011	026	
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OTHER SOURCE(S): MARPAT 138:255008

Epothilones, such as I [R0 = aryl, heteroaryl, arylalkyl, arylalkenyl, AΒ arylalkynyl, etc.; R1, R1', R2, R2' = H, alkyl, haloalkyl, etc.; R3, R3' = H, alkyl, etc.; R12 = H, OH, NH2, halogen, alkoxy, alkylamino, etc.; A-B, C-D = C(R1):C(R2), CR1R1'CR2R2', etc.; X = 0, S, CR3R3', NR3; Y = (CH2)m;Z = (CH2)q; m = 0-3, q = 1-3, and m + q = 1-4], were prepared for use in pharmaceutical compns. for the treatment of cancer. Thus, epothilone II was prepared via a multistep synthetic sequence which included an intramol. metathesis macrocyclization reaction using Grubbs' imidazole catalyst. The prepared epothilones were tested for cytotoxicity against a number of cancer cell lines.

IT 502619-65-0P

> RL: PAC (Pharmacological activity); PNU (Preparation, unclassified); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prepn of epothilones for therapeutic use as anticancer agents)

RN 502619-65-0 CAPLUS

4,17-Dioxabicyclo[14.1.0]heptadecane-5,9-dione, 7,11-dihydroxy-8,10,12,16-CN tetramethyl-3-[(1E)-1-methyl-2-(2-methyl-4-thiazolyl)ethenyl]-, (1S, 3S, 7S, 10R, 11S, 12S, 16R) - (9CI) (CA INDEX NAME)

Absolute stereochemistry. Double bond geometry as shown.

L6 ANSWER 5 OF 6 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2001:413810 CAPLUS

DOCUMENT NUMBER: 135:179755

TITLE: New Natural Epothilones from Sorangium cellulosum,

Strains So ce90/B2 and So ce90/D13: Isolation,

Structure Elucidation, and SAR Studies

AUTHOR(S): Hardt, Ingo H.; Steinmetz, Heinrich; Gerth, Klaus;

Sasse, F.; Reichenbach, Hans; Hoefle, Gerhard

CORPORATE SOURCE: Gesellschaft fuer Biotechnologische Forschung mbH,

Brown about D 20124 Common of the Polischia

Braunschweig, D-38124, Germany

SOURCE: Journal of Natural Products (2001), 64(7), 847-856

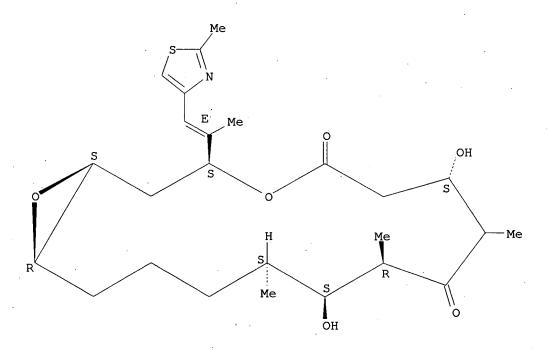
CODEN: JNPRDF; ISSN: 0163-3864

PUBLISHER: American Chemical Society

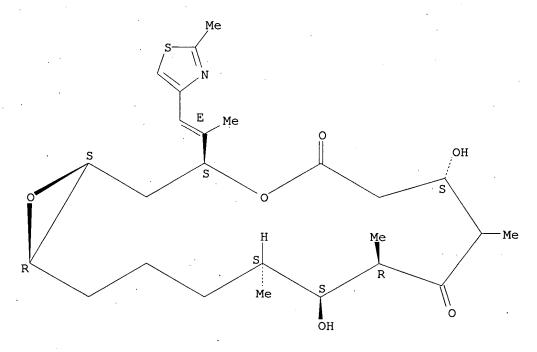
DOCUMENT TYPE: Journal LANGUAGE: English

In addition to epothilones A'(1) and B (2), 37 natural epothilone variants and epothilone-related compds. were isolated from the culture broth of a 700 L fermentation of Sorangium cellulosum, strain So ce90/B2. Of these, only the 12,13-desoxyepothilones, epothilone C (14) and D (15), were produced in significant amts. (3-6 mg/L); the 21-hydroxy derivs. and epothilones E (3) and F (4), in low and variable amts. due to further degradation by the producing organism. Most of the other epothilone variants were produced only in $1-100~\mu g/L$ amts. The new compds. are very similar in structure to the parent compds. 1, 2 and 14, 15 and are presumably the result of the imperfect selectivity of the biosynthetic enzymes for acetate and propionate. Further, epothilones containing an oxazole moiety (10-13) in the side chain instead of a thiazole as well as ring-expanded 18-membered macrolides, epothilones I (30-35), and a ring contracted 14-membered macrolide, epothilone K (36), were found as very minor metabolites. The mutant strain, So ce90/D13, instead of macrolactones, produced short-chain carboxylic acids 40, 41, and 42 bearing the characteristic thiazole side chain. The structures of the new epothilones were elucidated on the basis of comprehensive NMR and MS data. The new epothilone variants were tested in a cytotoxicity assay with mouse fibroblasts (cell line L929), and structure-activity relationships were established. Several new natural epothilones showed activity comparable to 1 and 2, but in no case exceeded that of 2.

Absolute stereochemistry. Rotation (-). Double bond geometry as shown. Currently available stereo shown.



RN 252917-30-9 CAPLUS
CN 4,17-Dioxabicyclo[14.1.0]heptadecane-5,9-dione, 7,11-dihydroxy-8,10,12-trimethyl-3-[(1E)-1-methyl-2-(2-methyl-4-thiazolyl)ethenyl]-,
(1S,3S,7S,10R,11S,12S,16R)- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 45 THERE ARE 45 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 6 OF 6 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

1999:811249 CAPLUS

DOCUMENT NUMBER:

132:49105

TITLE:

Epothilone minor constituents

INVENTOR(S):

Hoefle, Gerhard; Reichenbach, Hans; Gerth, Klaus;

Hardt, Ingo; Sasse, Florenz; Steinmetz, Heinrich

PATENT ASSIGNEE(S):

Gesellschaft Fur Biotechnologische Forschung m.b.H.

(Gbf), Germany

SOURCE:

PCT Int. Appl., 36 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

German

FAMILY ACC. NUM. COUNT:

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PATENT INFORMATION:

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CA	2336	189					1999									9990	
ΑU	9948	995					2000								•	9990	
ΑU	7574	52					2003						•		_	,,,,	010
EP	1087	975								EP 1	999_	9327	იი		1	9990	618
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JP 2002518397
                                 20020625
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                                             EP 2002-22332
                                                                     19990618
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                                 20040130
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                                             US 2001-719932
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                                 20060629
                                             US 2006-354769
                                                                     20060215
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                                 20070626
PRIORITY APPLN. INFO.:
                                             DE 1998-19826988
                                                                  A 19980618
                                             EP 1999-932700
                                                                  A3 19990618
                                             WO 1999-EP4244
                                                                  W
                                                                    19990618
                                             US 2001-719932
                                                                  A3 20010321
                                             US 2003-457098
                                                                  A1 20030606
```

AB The invention relates to compds. which are obtained by fermenting DSM 6773, especially epothilones A1, A2, A8, A9, B10, C1, C2, C3, C4, C5, C6, C7, C8, C9, D1, D2, D5, G1, G2, H1, H2, I1, I2, I3, I4, I5, I6 and K and trans-epothilones C1 and C2.

IT 252917-29-6P, Epothilone Al 252917-30-9P, Epothilone A2
RL: BAC (Biological activity or effector, except adverse); BOC (Biological occurrence); BSU (Biological study, unclassified); PRP (Properties); PUR (Purification or recovery); BIOL (Biological study); OCCU (Occurrence); PREP (Preparation)

(epothilone minor constituents)

RN 252917-29-6 CAPLUS

CN 4,17-Dioxabicyclo[14.1.0]heptadecane-5,9-dione, 7,11-dihydroxy-8,10,12-trimethyl-3-[(1E)-1-methyl-2-(2-methyl-4-thiazolyl)ethenyl]-, (1S,3S,7S,10R,11S,12S,16R)- (9CI) (CA INDEX NAME)

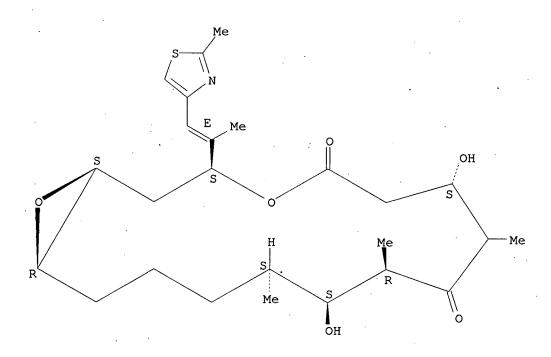
Absolute stereochemistry. Rotation (-). Double bond geometry as shown. Currently available stereo shown.

RN 252917-30-9 CAPLUS

CN 4,17-Dioxabicyclo[14.1.0]heptadecane-5,9-dione, 7,11-dihydroxy-8,10,12-

trimethyl-3-[(1E)-1-methyl-2-(2-methyl-4-thiazolyl)ethenyl]-, (1S, 3S, 7S, 10R, 11S, 12S, 16R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (+). Double bond geometry as shown. Currently available stereo shown.



=> d his

(FILE 'HOME' ENTERED AT 10:19:07 ON 13 AUG 2007)

FILE 'REGISTRY' ENTERED AT 10:19:20 ON 13 AUG 2007
L1 STRUCTURE UPLOADED
L2 STRUCTURE UPLOADED

L3 14 S L1 FULL L4 9 S L2 FULL

FILE 'CAPLUS' ENTERED AT 10:20:57 ON 13 AUG 2007

L5 3 S L3 FULL L6 6 S L4 FULL

CA SUBSCRIBER PRICE

=> log y
COST IN U.S. DOLLARS
SINCE FILE TOTAL
ENTRY SESSION
FULL ESTIMATED COST
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)
SINCE FILE TOTAL
ENTRY SESSION

-7.02

-7.02

STN INTERNATIONAL LOGOFF AT 10:22:40 ON 13 AUG 2007

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Welcome to STN International! Enter x:x

LOGINID: SSPTANXR1625

PASSWORD:

NEWS LOGIN

NEWS IPC8

TERMINAL (ENTER 1, 2, 3, OR ?):2

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Welcome to STN International
NEWS 1
                Web Page for STN Seminar Schedule - N. America
NEWS 2 MAY 01
                New CAS web site launched
NEWS
     3 MAY 08
                CA/CAplus Indian patent publication number format defined
NEWS
     4 MAY 14
                RDISCLOSURE on STN Easy enhanced with new search and display
                 fields
NEWS
        MAY 21
                BIOSIS reloaded and enhanced with archival data
     6 MAY 21
                TOXCENTER enhanced with BIOSIS reload
NEWS
     7 MAY 21
NEWS
                CA/CAplus enhanced with additional kind codes for German
                patents
        MAY 22
NEWS
     8
                CA/CAplus enhanced with IPC reclassification in Japanese
                patents
NEWS 9
        JUN 27
                CA/CAplus enhanced with pre-1967 CAS Registry Numbers
NEWS 10
        JUN 29
                STN Viewer now available
        JUN 29
NEWS 11
                STN Express, Version 8.2, now available
NEWS 12
        JUL 02
                LEMBASE coverage updated
NEWS 13
                LMEDLINE coverage updated
        JUL 02
NEWS 14
        JUL 02
                SCISEARCH enhanced with complete author names
        JUL 02
NEWS 15
                CHEMCATS accession numbers revised
NEWS 16
        JUL 02
                CA/CAplus enhanced with utility model patents from China
NEWS 17
        JUL 16 CAplus enhanced with French and German abstracts
NEWS 18 JUL 18 CA/CAplus patent coverage enhanced
NEWS 19 JUL 26 USPATFULL/USPAT2 enhanced with IPC reclassification
NEWS 20 JUL 30 USGENE now available on STN
NEWS 21 AUG 06 CAS REGISTRY enhanced with new experimental property tags
NEWS 22 AUG 06 BEILSTEIN updated with new compounds
NEWS 23 AUG 06 FSTA enhanced with new thesaurus edition
NEWS 24 AUG 13 CA/CAplus enhanced with additional kind codes for granted
                patents
NEWS EXPRESS
             29 JUNE 2007: CURRENT WINDOWS VERSION IS V8.2,
              CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
              AND CURRENT DISCOVER FILE IS DATED 05 JULY 2007.
NEWS HOURS
              STN Operating Hours Plus Help Desk Availability
```

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For general information regarding STN implementation of IPC 8

FILE 'HOME' ENTERED AT 10:24:33 ON 13 AUG 2007

=> FILE REG COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION

0.21

0.21

FULL ESTIMATED COST

FILE 'REGISTRY' ENTERED AT 10:25:05 ON 13 AUG 2007 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS.
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STRUCTURE FILE UPDATES: 12 AUG 2007 HIGHEST RN 944447-30-7 DICTIONARY FILE UPDATES: 12 AUG 2007 HIGHEST RN 944447-30-7

New CAS Information Use Policies, enter HELP USAGETERMS for details.

TSCA INFORMATION NOW CURRENT THROUGH December 2, 2006

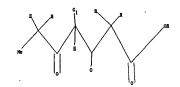
Please note that search-term pricing does apply when conducting SmartSELECT searches.

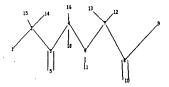
REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

http://www.cas.org/support/stngen/stndoc/properties.html

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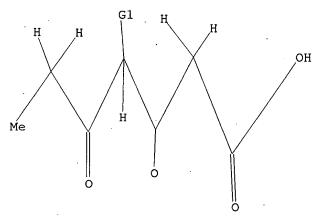


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chain nodes :
1  2  3  4  5  6  7  8  9  10  11  12  13  14  15  16  18
chain bonds :
1-2  2-3  2-14  2-15  3-4  3-5  4-6  4-16  4-18  6-7  6-11  7-8  7-12  7-13  8-9
8-10
exact/norm bonds :
3-5  4-16  6-11
exact bonds :
1-2  2-3  2-14  2-15  3-4  4-6  4-18  6-7  7-8  7-12  7-13
normalized bonds :
8-9  8-10
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G1:H,C

Match level:
1:CLASS 2:CLASS 3:CLASS 4:CLASS 5:CLASS 6:CLASS 7:CLASS 8:CLASS 9:CLASS 10:CLASS 11:CLASS 12:CLASS 13:CLASS 14:CLASS 15:CLASS 16:CLASS 18:CLASS

=> d 11 L1 HAS NO ANSWERS L1 STR



G1 H,C

Structure attributes must be viewed using STN Express query preparation.

=> s l1 full

FULL SEARCH INITIATED 10:25:50 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 113937 TO ITERATE

100.0% PROCESSED 113937 ITERATIONS

3 ANSWERS

SEARCH TIME: 00.00.02

L2 3 SEA SSS FUL L1

=> file caplus COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 172.10 172.31

COST IN U.S. DOLLARS
FULL ESTIMATED COST

FILE 'CAPLUS' ENTERED AT 10:25:57 ON 13 AUG 2007 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS.

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FILE COVERS 1907 - 13 Aug 2007 VOL 147 ISS 8 FILE LAST UPDATED: 12 Aug 2007 (20070812/ED)

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http://www.cas.org/infopolicy.html

=> s 13 full L3 NOT FOUND

The L-number entered has not been defined in this session, or it has been deleted. To see the L-numbers currently defined in this session, enter DISPLAY HISTORY at an arrow prompt (=>).

=> s 12 full L3 2 L2

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L3 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

2004:1068138 CAPLUS

DOCUMENT NUMBER:

142:197448

TITLE:

Highly Efficient Nickel-Catalyzed Cross-Coupling of Succinic and Glutaric Anhydrides with Organozinc

Reagents

AUTHOR(S):

Bercot, Eric A.; Rovis, Tomislav

CORPORATE SOURCE:

Department of Chemistry, Colorado State University,

Fort Collins, CO, 80523, USA

SOURCE:

Journal of the American Chemical Society (2005),

127(1), 247-254

CODEN: JACSAT; ISSN: 0002-7863

PUBLISHER:

American Chemical Society

DOCUMENT TYPE: LANGUAGE:

Journal English

OTHER SOURCE(S):

CASREACT 142:197448

AB A nickel-catalyzed alkylation of succinic and glutaric anhydrides with alkyl- and arylzinc reagents has been developed. A dramatic olefin effect has been investigated resulting in the identification of several styrene-based promoters which show pronounced enhancements in reaction rate. The substrate scope with respect to electrophilic and nucleophilic coupling partners has been examined and found to be remarkably broad, allowing for rapid introduction of mol. complexity through the use of functionalized coupling partners. Regioselective alkylation of an unsym. succinic anhydride and a profound effect of pendent coordinating olefins on reaction rate suggest a mechanism involving discrete oxidative addition of the nickel complex into the cyclic anhydride followed by a transmetalation event.

IT 838906-37-9P 838906-40-4P

RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (alkylation on nickel-catalyzed cross-coupling of succinic and glutaric anhydrides with organozinc reagents)

RN 838906-37-9 CAPLUS

CN Heptanoic acid, 5-oxo-3-(phenylmethoxy)- (9CI) (CA INDEX NAME)

RN 838906-40-4 CAPLUS

CN Heptanoic acid, 3-(acetyloxy)-3-methyl-5-oxo- (9CI) (CA INDEX NAME)

$$\begin{array}{c|c} \text{Me} & \text{O} \\ & | & || \\ \text{HO}_2\text{C}-\text{CH}_2-\text{C}-\text{CH}_2-\text{C}-\text{Et} \\ & | \\ & \text{OAc} \end{array}$$

REFERENCE COUNT:

82 THERE ARE 82 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

2004:550960 CAPLUS

DOCUMENT NUMBER:

141:106321

TITLE:

Preparation of epothilone derivatives for use in pharmaceutical compositions as antitumor agents Denni-Dischert, Donatienne; Floersheimer, Andreas;

INVENTOR(S):

Kuesters, Ernst; Oberer, Lukas; Sedelmeier, Gottfried

PATENT ASSIGNEE(S):

Novartis A.-G., Switz.; Novartis Pharma G.m.b.H.

SOURCE:

PCT Int. Appl., 50 pp.

•

CODEN: PIXXD2

DOCUMENT TYPE: LANGUAGE:

Patent English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

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	WO																	
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		•																
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			-		ZA,													
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			SI,	SK,	·TR							•						-
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		1732										2003-					0031	
	JР	2006	5140	2.5		T		2006	0427		JP.	2004-	-561 <i>4</i>	16		2		
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GT	. 50	OICE	(5).			1.11.71./1		T-4 T :	10032	<u> </u>								

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

AB C4-demethyl-epothilones or C4-bisnor-epothilones, such as I [R1, R7 = H, alkyl; R2 = nitrogen containing heteroaryl; R3 = H, Me; X = O, NR7; Z = O, bond], were prepared via fermentation and organic synthesis for use in pharmaceutical

compns. as antitumor agents. Thus, C4-bisnor-epothilone B II (R3 = H) was

prepared via an aldol condensation of aldehyde III with in situ disilylated (3R)-3-hydroxy-5-oxoheptanoic acid followed by a desilylation/macrolactonization reaction sequence. Also, C4-demethyl-epothilone B II (R = Me) was prepared directly by a fermentation process. The prepared epothilones were assayed for efficacy against human KB-31 and KB-8511 carcinoma cells. Drug delivery formulations containing the prepared epithilones were presented.

TT 717917-50-5, (3R)-3-Hydroxy-5-oxoheptanoic acid
RL: RCT (Reactant); RACT (Reactant or reagent)

(preparation of epothilone derivs. via fermentation and organic synthesis for use in

pharmaceutical compns. as antitumor agents)

RN 717917-50-5 CAPLUS

CN Heptanoic acid, 3-hydroxy-5-oxo-, (3R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

=> d his

(FILE 'HOME' ENTERED AT 10:24:33 ON 13 AUG 2007)

FILE 'REGISTRY' ENTERED AT 10:25:05 ON 13 AUG 2007

L1 STRUCTURE UPLOADED

L2 3 S L1 FULL

FILE 'CAPLUS' ENTERED AT 10:25:57 ON 13 AUG 2007 L3 2 S L2 FULL

=> log y

COST IN U.S. DOLLARS

SINCE FILE TOTAL
ENTRY SESSION
11.01 183.32

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE TOTAL

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Welcome to STN International! Enter x:x

LOGINID: SSPTANXR1625

PASSWORD:

NEWS HOURS

NEWS LOGIN

NEWS IPC8

TERMINAL (ENTER 1, 2, 3, OR ?):2

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NEWS
                 Web Page for STN Seminar Schedule - N. America
NEWS
                New CAS web site launched
        MAY 01
NEWS
     3
        MAY 08
                CA/CAplus Indian patent publication number format defined
NEWS 4
        MAY 14
                 RDISCLOSURE on STN Easy enhanced with new search and display
                 fields
                 BIOSIS reloaded and enhanced with archival data
NEWS 5
        MAY 21
NEWS
                TOXCENTER enhanced with BIOSIS reload
NEWS
        MAY 21
                CA/CAplus enhanced with additional kind codes for German
                 patents
NEWS
        MAY 22
                CA/CAplus enhanced with IPC reclassification in Japanese
                 patents
         JUN 27
NEWS 9
                CA/CAplus enhanced with pre-1967 CAS Registry Numbers
NEWS 10
         JUN 29
                STN Viewer now available
         JUN 29
NEWS 11
                 STN Express, Version 8.2, now available
NEWS 12
         JUL 02
                LEMBASE coverage updated
NEWS 13
         JUL 02
                LMEDLINE coverage updated
NEWS 14
         JUL 02
                 SCISEARCH enhanced with complete author names
NEWS 15
         JUL 02
                 CHEMCATS accession numbers revised
NEWS 16
         JUL 02
                CA/CAplus enhanced with utility model patents from China
NEWS 17
         JUL 16
                CAplus enhanced with French and German abstracts
NEWS 18
         JUL 18
                CA/CAplus patent coverage enhanced
NEWS 19
         JUL 26
                USPATFULL/USPAT2 enhanced with IPC reclassification
NEWS 20
        JUL 30
                USGENE now available on STN
NEWS 21 AUG 06 CAS REGISTRY enhanced with new experimental property tags
NEWS 22 AUG 06 BEILSTEIN updated with new compounds
NEWS 23 AUG 06 FSTA enhanced with new thesaurus edition
NEWS 24
       AUG 13 CA/CAplus enhanced with additional kind codes for granted
                 patents
NEWS EXPRESS 29 JUNE 2007: CURRENT WINDOWS VERSION IS V8.2,
              CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
              AND CURRENT DISCOVER FILE IS DATED 05 JULY 2007.
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FILE 'HOME' ENTERED AT 10:27:04 ON 13 AUG 2007

=> FILE REG COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 0.21 0.21

FULL ESTIMATED COST

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STRUCTURE FILE UPDATES: 12 AUG 2007 HIGHEST RN 944447-30-7 DICTIONARY FILE UPDATES: 12 AUG 2007 HIGHEST RN 944447-30-7

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http://www.cas.org/support/stngen/stndoc/properties.html

=>

Uploading C:\Program Files\Stnexp\Queries\10538200g.str

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exact/norm bonds:
6-10 11-13 14-16 22-23 25-26
exact bonds:
2-5 5-6 5-7 5-8 6-9 9-11 9-30 11-12 12-14 12-15 14-17 14-31 17-18
17-19 17-32 19-20 20-21 21-22 23-24 24-25 25-27 27-28 27-29
normalized bonds:
2-3 2-4

G1:H,C

=>

Match level:
2:CLASS 3:CLASS 4:CLASS 5:CLASS 6:CLASS 7:CLASS 8:CLASS 9:CLASS 10:CLASS
11:CLASS 12:CLASS 13:CLASS 14:CLASS 15:CLASS 16:CLASS 17:CLASS 18:CLASS
19:CLASS 20:CLASS 21:CLASS 22:CLASS 23:CLASS 24:CLASS 25:CLASS 26:CLASS 27:CLASS 28:CLASS 29:CLASS 30:CLASS 31:CLASS 32:CLASS

L1 STRUCTURE UPLOADED

Uploading C:\Program Files\Stnexp\Queries\10538200c.str

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2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 24 25 26
27 28 29 30 31 32
ring nodes :
22 23 33
chain bonds :
2-3 2-4 2-5 5-6 5-7 5-8 6-9 6-10 9-11 9-30 11-12 11-13 12-14 12-15
14-16 14-17 14-31 17-18 17-19 17-32 19-20 20-21 21-22 23-24 24-25 25-26
25-27 27-28 27-29
ring bonds :
22-23 22-33 23-33
exact/norm bonds :
6-10 11-13 14-16 22-23 22-33 23-33 25-26
exact bonds :

2-5 5-6 5-7 5-8 6-9 9-11 9-30 11-12 12-14 12-15 14-17 14-31 17-18 17-19 17-32 19-20 20-21 21-22 23-24 24-25 25-27 27-28 27-29 normalized bonds : 2-3 2-4

G1:H,C

Match level:

2:CLASS 3:CLASS 4:CLASS 5:CLASS 6:CLASS 7:CLASS 8:CLASS 9:CLASS 10:CLASS 11:CLASS 12:CLASS 13:CLASS 14:CLASS 15:CLASS 16:CLASS 17:CLASS 18:CLASS 19:CLASS 20:CLASS 21:CLASS 22:CLASS 23:CLASS 24:CLASS 25:CLASS 26:CLASS 27:CLASS 28:CLASS 29:CLASS 30:CLASS 31:CLASS 32:CLASS 33:Atom

L2 STRUCTURE UPLOADED

=> d 11 L1 HAS NO ANSWERS L1 STR

Structure attributes must be viewed using STN Express query preparation.

=> s 11 full
FULL SEARCH INITIATED 10:28:31 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 912 TO ITERATE

100.0% PROCESSED 912 ITERATIONS SEARCH TIME: 00.00.01

0 ANSWERS

L3

G1 H,C

0 SEA SSS FUL L1

=> d 12 L2 HAS NO ANSWERS L2 STR

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT * Structure attributes must be viewed using STN Express query preparation.

=> s 12 full FULL SEARCH INITIATED 10:28:47 FILE 'REGISTRY' FULL SCREEN SEARCH COMPLETED - 169 TO ITERATE

100.0% PROCESSED 169 ITERATIONS

0 ANSWERS

SEARCH TIME: 00.00.01

ή.4

0 SEA SSS FUL L2

=> log y

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

2.4

ENTRY SESSION

FULL ESTIMATED COST

344.20 344.41

STN INTERNATIONAL LOGOFF AT 10:28:51 ON 13 AUG 2007

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PASSWORD:

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                 Web Page for STN Seminar Schedule - N. America
NEWS
      1
NEWS
      2 MAY 01
                 New CAS web site launched
NEWS
      3 MAY 08
                 CA/CAplus Indian patent publication number format defined
NEWS
        MAY 14
                 RDISCLOSURE on STN Easy enhanced with new search and display
                 fields
NEWS
         MAY 21
                 BIOSIS reloaded and enhanced with archival data
NEWS
        MAY 21
                 TOXCENTER enhanced with BIOSIS reload
NEWS
        MAY 21
                 CA/CAplus enhanced with additional kind codes for German
                 patents
         MAY 22
NEWS
      8
                 CA/CAplus enhanced with IPC reclassification in Japanese
                 patents
NEWS
         JUN 27
                 CA/CAplus enhanced with pre-1967 CAS Registry Numbers
NEWS 10
         JUN 29
                 STN Viewer now available
         JUN 29
NEWS 11
                 STN Express, Version 8.2, now available
NEWS 12
         JUL 02
                 LEMBASE coverage updated
NEWS 13
         JUL 02
                 LMEDLINE coverage updated
NEWS 14
         JUL 02
                 SCISEARCH enhanced with complete author names
         JUL 02
NEWS 15
                 CHEMCATS accession numbers revised
NEWS 16
         JUL 02
                 CA/CAplus enhanced with utility model patents from China
NEWS 17
         JUL 16
                 CAplus enhanced with French and German abstracts
NEWS 18
         JUL 18
                CA/CAplus patent coverage enhanced
NEWS 19
         JUL 26
                USPATFULL/USPAT2 enhanced with IPC reclassification
        JUL 30
NEWS 20
                USGENE now available on STN
        AUG 06 CAS REGISTRY enhanced with new experimental property tags
NEWS 21
        AUG 06 BEILSTEIN updated with new compounds
NEWS 22
NEWS 23
         AUG 06
                 FSTA enhanced with new thesaurus edition
                 CA/CAplus enhanced with additional kind codes for granted
NEWS 24
        AUG 13
                 patents
NEWS EXPRESS
              29 JUNE 2007: CURRENT WINDOWS VERSION IS V8.2,
              CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
              AND CURRENT DISCOVER FILE IS DATED 05 JULY 2007.
              STN Operating Hours Plus Help Desk Availability
NEWS HOURS
NEWS LOGIN
              Welcome Banner and News Items
NEWS IPC8
              For general information regarding STN implementation of IPC 8
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Enter NEWS followed by the item number or name to see news on that specific topic.

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=> FILE REG

COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 0.21 0.21

FULL ESTIMATED COST

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Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

12 AUG 2007 HIGHEST RN 944447-30-7 STRUCTURE FILE UPDATES: DICTIONARY FILE UPDATES: 12 AUG 2007 HIGHEST RN 944447-30-7

New CAS Information Use Policies, enter HELP USAGETERMS for details.

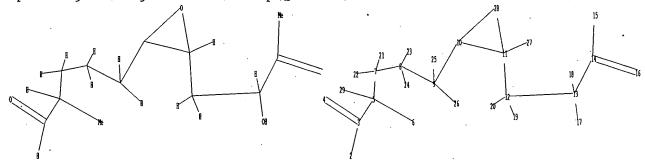
TSCA INFORMATION NOW CURRENT THROUGH December 2, 2006

Please note that search-term pricing does apply when conducting SmartSELECT searches.

REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

http://www.cas.org/support/stngen/stndoc/properties.html

Uploading C:\Program Files\Stnexp\Queries\10538200e.str



chain nodes : 8 9 12 13 14

15 16 17 18 19 20 21 22 27 29

ring nodes : 10 11 28

chain bonds :

2-3 3-4 3-5 5-6 5-7 5-29 7-8 7-21 7-22 8-9 8-23 8-24 9-10 9-25 9-26

11-12 11-27 12-13 12-19 12-20 13-14 13-17 13-18 14-15 14-16

ring bonds:

10-11 10-28 11-28 exact/norm bonds :

3-4 10-11 10-28 11-28 13-17

exact bonds:
2-3 3-5 5-6 5-7 5-29 7-8 7-21 7-22 8-9 8-23 8-24 9-10 9-25 9-26 11-12 11-27 12-13 12-19 12-20 13-14 13-18 14-15 14-16

G1:H,C

Match level :

2:CLASS 3:CLASS 4:CLASS 5:CLASS 6:CLASS 7:CLASS 8:CLASS 9:CLASS 10:Atom 11:Atom 12:CLASS 13:CLASS 14:CLASS 15:CLASS 16:CLASS 17:CLASS 18:CLASS 19:CLASS 20:CLASS 21:CLASS 22:CLASS 23:CLASS 24:CLASS 25:CLASS 26:CLASS 27:CLASS 28:Atom 29:CLASS

L1 STRUCTURE UPLOADED

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Uploading C:\Program Files\Stnexp\Queries\10538200f.str

chain nodes:
2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23 24
25 26 27 28
chain bonds:
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10-11 11-12 11-27 12-13 12-19 12-20 13-14 13-17 13-18 14-15 14-16
exact/norm bonds:
3-4 10-11 13-17
exact bonds:
2-3 3-5 5-6 5-7 5-28 7-8 7-21 7-22 8-9 8-23 8-24 9-10 9-25 9-26 11-12
11-27 12-13 12-19 12-20 13-14 13-18 14-15 14-16

G1:H,C

Match level :

2:CLASS 3:CLASS 4:CLASS 5:CLASS 6:CLASS 7:CLASS 8:CLASS 9:CLASS 10:Atom 11:Atom 12:CLASS 13:CLASS 14:CLASS 15:CLASS 16:CLASS 17:CLASS 18:CLASS 19:CLASS 20:CLASS 21:CLASS 22:CLASS 23:CLASS 24:CLASS 25:CLASS 26:CLASS 27:CLASS 28:CLASS

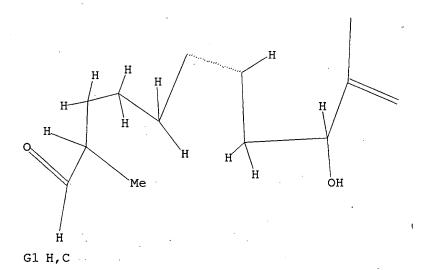
STRUCTURE UPLOADED

=> D L1

L2

STR

Structure attributes must be viewed using STN Express query preparation.



Structure attributes must be viewed using STN Express query preparation.

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FULL SEARCH INITIATED 10:31:16 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 1531 TO ITERATE

100.0% PROCESSED 1531 ITERATIONS SEARCH TIME: 00.00.01

2 ANSWERS

L3

2 SEA SSS FUL L1

=> s 12 full

FULL SEARCH INITIATED 10:31:20 FILE 'REGISTRY' FULL SCREEN SEARCH COMPLETED -

100.0% PROCESSED 69139 ITERATIONS

SEARCH TIME: 00.00.01

0 ANSWERS

L4

O SEA SSS FUL L2

=> file caplus

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

ENTRY

SESSION

344.20

344.41

FULL ESTIMATED COST

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=> s 13 full

1 L3

=> d ibib abs hitstr tot

ANSWER 1 OF 1 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

2004:550960 CAPLUS

DOCUMENT NUMBER:

TITLE:

Preparation of epothilone derivatives for use in pharmaceutical compositions as antitumor agents

INVENTOR(S):

Denni-Dischert, Donatienne; Floersheimer, Andreas; Kuesters, Ernst; Oberer, Lukas; Sedelmeier, Gottfried Novartis A.-G., Switz.; Novartis Pharma G.m.b.H.

PATENT ASSIGNEE(S): SOURCE:

PCT Int. Appl., 50 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004056832	A2	20040708	WO 2003-EP14747	20031222
WO 2004056832	A3	20040910		

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PRIORITY APPLN. INFO .:
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                                                                     W 20031222
                                               WO 2003-EP14747
OTHER SOURCE(S):
                          MARPAT 141:106321
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* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

AB C4-demethyl-epothilones or C4-bisnor-epothilones, such as I [R1, R7 = H, alkyl; R2 = nitrogen containing heteroaryl; R3 = H, Me; X = O, NR7; Z = O, bond], were prepared via fermentation and organic synthesis for use in pharmaceutical

compns. as antitumor agents. Thus, C4-bisnor-epothilone B II (R3 = H) was prepared via an aldol condensation of aldehyde III with in situ disilylated (3R)-3-hydroxy-5-oxoheptanoic acid followed by a desilylation/macrolactonization reaction sequence. Also, C4-demethyl-epothilone B II (R = Me) was prepared directly by a fermentation process. The prepared epothilones were assayed for efficacy against human KB-31 and KB-8511 carcinoma cells. Drug delivery formulations containing the prepared epithilones were presented.

IT 717917-44-7P

RL: BPN (Biosynthetic preparation); RCT (Reactant); BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent)

(preparation of epothilone derivs. via fermentation and organic synthesis for use in

pharmaceutical compns. as antitumor agents)

RN 717917-44-7 CAPLUS

CN Oxiranepentanal, $3-[(2S,3E)-2-hydroxy-3-methyl-4-(2-methyl-4-thiazolyl)-3-butenyl]-\alpha, 2-dimethyl-, (<math>\alpha S,2R,3S$)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.
Double bond geometry as shown.

Me
$$\sim$$
 N \sim N \sim

IT 717917-46-9P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU.

(Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

 $% \left(A_{i}\right) =A_{i}\left(A_{i}\right) +A_{i}\left(A_{i}\right) +A_{i}\left($

pharmaceutical compns. as antitumor agents)

RN 717917-46-9 CAPLUS

CN Oxiranepentanal, $3-[(2S,3E)-2-hydroxy-3-methyl-4-(2-methyl-4-thiazolyl)-3-butenyl]-\alpha-methyl-, (<math>\alpha S,2R,3S$)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.
Double bond geometry as shown.

=> d his

(FILE 'HOME' ENTERED AT 10:29:42 ON 13 AUG 2007)

FILE 'REGISTRY' ENTERED AT 10:30:04 ON 13 AUG 2007

L1 STRUCTURE UPLOADED

L2 STRUCTURE UPLOADED

L3 2 S L1 FULL L4 0 S L2 FULL

FILE 'CAPLUS' ENTERED AT 10:31:25 ON 13 AUG 2007 L5 1 S L3 FULL

=> log y

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FULL ESTIMATED COST ENTRY SESSION 5.74 350.15

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) SINCE FILE TOTAL

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